

## ADDITIVES FOR PAINTS AND INKS

[Technical field to which the invention belongs]

5                This invention relates to additives for paints and inks. The additives of the present invention can impart, when added to paints or inks, to the paint-coated or ink-printed surfaces, defoaming and anti-popping property or flow-and-leveling property. The additives of the invention furthermore have a property to prevent whitening  
10 phenomenon of painted surfaces and, therefore, is particularly useful as additives for clear paints for which transparency is the most important requirement. Whitening phenomenon of painted surfaces refers to a phenomenon of white blooming or turbidity caused in painted surfaces, due to infiltration of water thereinto. Hereinafter  
15 the property or ability to prevent this phenomenon is referred to as anti-whitening property. This anti-whitening property is evaluated by visual examination of extent of whitening caused after immersion of dry painted films in warm water.

20 [Prior art]

              Paints are generally blended with defoamers (defoaming and anti-popping agent) for removing foams which are entrained during paint application, or with flow-and-leveling agents for improving wettability of painting objects with paints and assisting level  
25 painting. When the defoamers and flow-and-leveling agents (which may hereafter be collectively referred to as surface control agents) are blended with paint, they remain in cured clear paint film in the form of particles. Some of the new curing type thermosetting or baking clear systems in which no melamine resin is used as the curing  
30 agent, which have recently appeared in the market as a countermeasure to acid precipitation, exhibit the whitening phenomenon due to

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In cured films of such new curing type paints, those blended surface control agents remain as fine particles. In consequence, when water penetrates into the interfaces between the surface control agent particles and the paint film, bubbles are formed in the surface control agent particles after the water is removed from the paint film, to impart the paint film white, turbid appearance. This whitening phenomenon is apt to impair appearance of the paints finish. On the other hand, reduction in the blended amount of surface control agents to avoid deterioration in clarity of the paint film often results in unsatisfactory exhibition of the primary functions of the

surface control agents.

Accordingly, therefore, the object of the present invention is to provide novel surface control agents for paints and inks, in particular, such surface control agents which do not cause whitening phenomenon and hence do not impair clarity of paint film, when they  
5 are blended in paints using no melamine resin as curing agent.

[Means to solve the problem]

We have engaged in extensive studies to discover that  
10 copolymers obtained by reacting reactive monomers used in conventional surface control agents with reactive monomers having isocyanate groups or those having groups derived from isocyanate groups can prevent whitening phenomenon without impairing the effects of the original surface control agents, and hence are useful as additives  
15 for, in particular, clear paint.

Thus, according to the present invention, a defoaming and anti-popping agent or flow-and-leveling agent for paints and inks is provided, which is characterized by comprising a copolymer obtained through copolymerization of 2-50% by weight of a reactive monomer  
20 having isocyanate group or an isocyanate-derived group with 98-50% by weight of other monomer or polymer which is reactable with said reactive monomer.

When the ratio of said monomer having isocyanate group or an isocyanate-derived group in the copolymer according to the  
25 invention is less than 2% by weight, the intended effect for preventing whitening phenomenon cannot be satisfactorily accomplished. On the other hand, when the ratio exceeds 50% by weight, the effects as a surface control agent are not fully exhibited or paint film properties are adversely affected.

30 Examples of reactive monomers having isocyanate groups include 2-isocyanatoethyl methacrylate, 2-isocyanatoethyl acrylate

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and 3-isopropenyl- $\alpha,\alpha$ -dimethylbenzyl isocyanate.

Examples of reactive monomers having isocyanate-derived groups include 2-(0-[1'-methylpropylideneamino]carboxy-amino)ethyl methacrylate and 2-(0-[1'-methylpropylideneamino]carboxy-amino)-  
5 ethyl acrylate.

As other monomers or polymers reactable with reactive monomers having isocyanate groups or isocyanate-derived groups, for example, alkyl esters of acrylic acid such as ethyl acrylate, butyl acrylate, 2-ethylhexyl acrylate and lauryl acrylate; alkyl esters of  
10 methacrylic acid such as ethyl methacrylate, butyl methacrylate, 2-ethylhexyl methacrylate, lauryl methacrylate, hexadecyl methacrylate and octadecyl methacrylate; alkyl vinyl ethers such as ethyl vinyl ether, propyl vinyl ether, butyl vinyl ether, 2-ethylhexyl vinyl ether, lauryl vinyl ether and octadecyl vinyl ether; reactive silicone having  
15 methacryloyloxy groups (Silaplane<sup>TM</sup> FM-0711 marketed by Chisso Corporation, AK-5 and AK-30 marketed by Toagosei Co., Ltd.); reactive acrylic polymers which are normally referred to as macromonomers (Diamac Series marketed by Mitsubishi Chemical Corporation and A-Series marketed by Toagosei Co., Ltd.); and polymers having  
20 reactive groups such as butadiene polymers can be named. Besides these, there are many kinds of monomers and polymers which are useful as starting materials of surface control agents, all of which are utilizable so long as they are copolymerizable with reactive monomers having isocyanate groups or isocyanate-derived groups.

25 Synthesis of a copolymer from above reactive monomers can be performed following radical polymerization process using peroxide or azo compound, cationic polymerization process using acid catalyst, or anionic polymerization process using alkali metal catalyst. The present invention concerns development of utility of the copoly-  
30 mers, and is in no way limited by their method of synthesis.

The surface control agents according to the present inven-

tion are usable with particular effectiveness for clear paints for which transparency of the finish is of predominant importance, e.g., clear top coat paint for automobiles in which resistance to whitening is a significant factor, and precoat metal (PCM) paint.

5           The surface control agents according to the present inven-  
tion can be added to paints and inks either during the process of  
preparing, or after the preparation, of the paints and inks.

The use rate of a surface control agent according to the invention is dependent on various factors such as nature of the paint or ink, conditions of use of the paint or ink, painting conditions of the paint or printing conditions of the ink and the like and hence no certain limits can be provided. Whereas, it is generally used within a range of 0.001 to 5.0%, preferably 0.01 to 2.0%, as converted to solid, based on the weight of the paint or ink. In case of its addition to paint, it may be blended in a greater amount than that of a conventional surface control agent for paint, because the former is less tended to cause whitening phenomenon.

[Effects of the invention]

20           The surface control agent for paint or ink according to the present invention is characterized by comprising a copolymer containing isocyanate groups or isocyanate-derived groups, and it not only imparts flow-and-leveling property or defoaming and anti-popping property to painted or printed surface but also prevents the phenomenon of paint film appearing white and turbid when it is added to, in  
25           particular, paint, even in the presence of particles of the surface control agent in the cured paint film.

### [Examples]

30 Hereinafter the invention is explained in further details,  
referring to Examples in which parts and percentages are by weight.

### Production Example 1 of a defoaming agent

A 1000 ml-reaction vessel equipped with a stirrer, reflux condenser, dropping funnel, thermometer and gaseous nitrogen inlet port was charged with 100 parts of xylene, which was heated to 100°C under introduction of gaseous nitrogen. Thereafter the following solution (a-1) was dropped into said xylene at a constant rate through the dropping funnel, consuming 90 minutes.

#### Solution (a-1)

	Octadecyl methacrylate	285 parts
10	2-Isocyanatoethyl methacrylate	15 parts
	Xylene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	5 parts

An hour after completion of the dropwise addition of the solution (a-1), 1.5 parts of the t-butylperoxy-2-ethyl hexanoate was added to the reaction system, which was allowed to react for subsequent 3 hours while its temperature was maintained at 100°C. After termination of the reaction, the solid content was adjusted to 30% using xylene, and a defoaming agent [DG-1] was obtained. So synthesized polymer had a weight-average molecular weight of 17,000.

### 20 Production Example 2 of a defoaming agent

A defoaming agent was prepared in the identical manner with above Production Example 1, except that the following solution (a-2) was added dropwise in place of the solution (a-1) used in Production Example 1.

#### 25 Solution (a-2)

	Hexadecyl methacrylate	190 parts
	Lauryl vinyl ether	80 parts
	2-(0-[1'-methylpropylideneamino]-carboxyamino)ethyl methacrylate	30 parts
30	Xylene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	6 parts

After termination of the reaction, the solid content of the reaction product was adjusted to 30% using xylene, and a defoaming agent [DG-2] was obtained. So synthesized polymer had a weight-average molecular weight of 25,000.

5 Production Example 3 of a defoaming agent

A defoaming agent was prepared in the identical manner with Production Example 1, except that the following solution (a-3) was added dropwise in place of the solution (a-1) used in Production Example 1.

10 Solution (a-3)

Lauryl methacrylate	155 parts
2-(0-[1'-methylpropylideneamino]- carboxyamino)ethyl methacrylate	145 parts
Xylene	100 parts
t-Butylperoxy-2-ethyl hexanoate	15 parts

After termination of the reaction, the solid content of the reaction product was adjusted to 30% with xylene, and a defoaming agent [DG-3] was obtained. So synthesized polymer had a weight-average molecular weight of 23,000.

20 Production Example 1 of a flow-and-leveling agent

A 1000 ml-reaction vessel equipped with a stirrer, reflux condenser, dropping funnel, thermometer and gaseous nitrogen inlet port was charged with 150 parts of toluene which was heated to 110°C and refluxed under introduction of gaseous nitrogen. Then the following solution (b-1) was dropped into the system at a constant rate through the dropping funnel, consuming 2 hours.

Solution (b-1)

2-Ethylhexyl acrylate	285 parts
2-Isocyanatoethyl methacrylate	15 parts
Toluene	100 parts
t-Butylperoxy-2-ethyl hexanoate	9 parts

An hour after completion of the dropwise addition of the solution (b-1), 3 parts of t-butylperoxy-2-ethylhexanoate was added and the system was allowed to react for subsequent 2 hours while its temperature was maintained at 110°C. After termination of the  
 5 reaction, the solid content of the reaction product was adjusted to 50% with toluene, and a flow-and-leveling agent [SG-1] was obtained. So synthesized polymer had a number-average molecular weight of 10,000.

Production Example 2 of a flow-and-leveling agent

10 The reaction was conducted in the identical manner with above Production Example 1 of the flow-and-leveling agent, except that the following solution (b-2) was used in place of the solution (b-1).

Solution (b-2)

	n-Butyl acrylate	190 parts
15	Isobutyl vinyl ether	80 parts
	2-(0-[1'-methylpropylideneamino]-carboxyamino)ethyl methacrylate	30 parts
	Toluene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	15 parts

20 After termination of the reaction, the solid content of the reaction product was adjusted to 50% using toluene, and a flow-and-leveling agent [SG-2] was obtained. So synthesized polymer had a number-average molecular weight of 3,500.

Production Example 3 of a flow-and-leveling agent

25 The reaction was conducted in the identical manner with the Production Example 1 of the flow-and-leveling agent, except that the following solution (b-3) was used in place of the solution (b-1).

Solution (b-3)

	n-Butyl acrylate	95 parts
30	Methacryloyloxypropylpolydimethylsiloxane	60 parts

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2-(0-[1'-methylpropylideneamino]-	
carboxyamino)ethyl methacrylate	145 parts
Toluene	100 parts
t-Butylperoxy-2-ethyl hexanoate	6 parts

5 After termination of the reaction, the solid content of the reaction product was adjusted to 50% using toluene, and a flow-and-leveling agent [SG-3] was obtained. So synthesized polymer had a number-average molecular weight of 9,000.

Comparative Example 1 of producing a defoaming agent

10 A reaction vessel similar to the one used in Production Example 1 of the defoaming agent was charged with 100 parts of xylene, which was heated to 100°C under introduction of gaseous nitrogen. Then the following solution (c-1) was added dropwise through the dropping funnel at a constant rate, consuming 90 min-  
15 utes.

Solution (c-1)

Octadecyl methacrylate	300 parts
Xylene	100 parts
t-Butylperoxy-2-ethyl hexanoate	5 parts

20 An hour after completion of the dropwise addition of the solution (c-1), 1.5 parts of t-butylperoxy-2-ethyl hexanoate was added, and the system was allowed to react for subsequent 3 hours while its temperature was maintained at 100°C. After termination of the reaction, the solid content of the reaction product was adjusted to  
25 30% using xylene, and a defoaming agent [DN-1] was obtained. So synthesized polymer had a weight-average molecular weight of 20,000.

Comparative Example 2 of producing a defoaming agent

The reaction of preceding Comparative Example 1 was repeated except that the following solution (c-2) was used in place of  
30 the solution (c-1).

Solution (c-2)

	Hexadecyl methacrylate	210 parts
	Lauryl vinyl ether	90 parts
	Xylene	100 parts
5	t-Butylperoxy-2-ethyl hexanoate	15 parts

After termination of the reaction, the solid content of the reaction product was adjusted to 30% with xylene, to provide a de-foaming agent [DN-2]. So synthesized polymer had a weight-average molecular weight of 30,000.

10 Comparative Example 3 of producing a defoaming agent

The reaction of Example 1 of producing the defoaming agent was repeated, except that the 2-isocyanatoethyl methacrylate used in the solution (a-1) of said Example 1 was replaced by 2-hydroxyethyl methacrylate (solution (c-3)).

15 Solution (c-3)

	Octadecyl methacrylate	285 parts
	2-Hydroxyethyl methacrylate	15 parts
	Xylene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	15 parts

20 After termination of the reaction, the solid content of the reaction product was adjusted to 30% with xylene, to provide a de-foaming agent [DN-3]. The synthesized polymer had a weight-average molecular weight of 24,000.

Comparative Example 4 of producing a defoaming agent

25 The reaction of Example 1 of producing the defoaming agent was repeated, except that the following solution (c-4) was used in place of the solution (a-1) in said Example 1.

Solution (c-4)

	Octadecyl methacrylate	295 parts
30	2-Isocyanatoethyl methacrylate	5 parts
	Xylene	100 parts

t-Butylperoxy-2-ethyl hexanoate	15 parts
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An hour after completion of the dropwise addition of the solution (c-4), 1.5 parts of the t-butylperoxy-2-ethyl hexanoate was added to the reaction system, which was allowed to react for subsequent 3 hours while its temperature was maintained at 100°C. After termination of the reaction, the solid content of the reaction product was adjusted to 30%, to provide a defoaming agent [DN-4]. The synthesized polymer had a weight-average molecular weight of 19,000.

### Comparative Example 5 of producing a defoaming agent

10           The reaction of Example 1 of producing the defoaming agent was repeated, except that the following solution (c-5) was used in place of the solution (a-1) in said Example 1.

Solution (c-5)

15	Octadecyl methacrylate	140 parts
	2-(0-[1'-methylpropylideneamino]- carboxyamino)ethyl methacrylate	160 parts
	Xylene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	15 parts

An hour after completion of the dropwise addition of the solution (c-5), 1.5 parts of the t-butylperoxy-2-ethyl hexanoate was added to the reaction system, which was allowed to react for subsequent 3 hours while its temperature was maintained at 100°C. After termination of the reaction, the solid content of the reaction product was adjusted to 30%, to provide a defoaming agent [DN-5]. The synthesized polymer had a weight-average molecular weight of 22,000.

### Comparative Example 1 of producing a flow-and-leveling agent

A reaction vessel similar to the one used in Production Example 1 of the flow-and-leveling agent was charged with 150 parts of toluene which was subsequently heated to 110°C and refluxed under introduction of gaseous nitrogen. Then the following solution (d-1) was dropped into said system at a constant rate through the

dropping funnel, consuming 2 hours.

Solution (d-1)

	2-Ethylhexyl acrylate	300 parts
	Toluene	100 parts
5	t-Butylperoxy-2-ethyl hexanoate	6 parts

An hour after completion of the dropwise addition of the solution (d-1), 3 parts of t-butylperoxy-2-ethylhexanoate was added and the system was allowed to react for subsequent 2 hours while its temperature was maintained at 110°C. After termination of the  
 10 reaction, the solid content of the reaction product was adjusted to 50% with toluene, and a flow-and-leveling agent [SN-1] was obtained. The synthesized polymer had a number-average molecular weight of 9,500.

Comparative Example 2 of producing a flow-and-leveling agent

15 The reaction was conducted in the identical manner with above Comparative Example 1 of producing the flow-and-leveling agent, except that the following solution (d-2) was used in place of the solution (d-1).

Solution (d-2)

20	n-Butyl acrylate	205 parts
	Isobutyl vinyl ether	95 parts
	Toluene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	15 parts

After termination of the reaction, the solid content of the  
 25 reaction product was adjusted to 50% using toluene, and a flow-and-leveling agent [SN-2] was obtained. The synthesized polymer had a number-average molecular weight of 3,000.

Comparative Example 3 of producing a flow-and-leveling agent

The reaction was conducted in the identical manner with  
 30 the Production Example 1 of the flow-and-leveling agent, except that the 2-isocyanatoethyl methacrylate in the solution (b-1) used in said

Example was replaced by N,N-dimethylacrylamide (the following solution (d-3)).

Solution (d-3)

	2-Ethylhexyl acrylate	270 parts
5	N,N-dimethylacrylamide	30 parts
	Toluene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	6 parts

After termination of the reaction, solid content of the reaction product was adjusted to 50% using toluene, and a flow-and-leveling agent [SN-3] was obtained. The synthesized polymer had a number-average molecular weight of 10,500.

Comparative Example 4 of producing a flow-and-leveling agent

The reaction of Production Example 2 of the flow-and-leveling agent was repeated except that the following solution (d-4) was used in place of the solution (b-2).

Solution (d-4)

	n-Butyl acrylate	205 parts
	Isobutyl vinyl ether	90 parts
20	2-(0-[1'-methylpropylideneamino]-carboxyamino)ethyl methacrylate	5 parts
	Toluene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	15 parts

After termination of the reaction, the solid content of the reaction product was adjusted to 50% with toluene, to provide a flow-and-leveling agent [SN-4]. The number-average molecular weight of the synthesized polymer was 3,500.

Comparative Example 5 of producing a flow-and-leveling agent

The reaction of Production Example 2 of the flow-and-leveling agent was repeated except that the following solution (d-5) was used in place of the solution (b-2).

Solution (d-5)

	n-Butyl acrylate	105 parts
	Isobutyl vinyl ether	35 parts
5	2-(0-[1'-methylpropylideneamino]- carboxyamino)ethyl methacrylate	160 parts
	Toluene	100 parts
	t-Butylperoxy-2-ethyl hexanoate	15 parts

After termination of the reaction, the solid content of the reaction product was adjusted to 50% using toluene, to provide a flow-  
 10 and-leveling agent [SN-5]. The synthesized polymer had a number-average molecular weight of 4,000.

TABLE 1

15 Additives formed in Production Examples of the defoaming agents

	Additive	Weight-average molecular weight	Non-volatile component (%)
20	Production Example 1	17000	30
	Production Example 2	25000	30
25	Production Example 3	23000	30

TABLE 2

Additives formed in Production Examples of the flow-and-leveling agents

5		Additive	Number-average molecular weight	Non-volatile component (%)
	Production Example 1	SG-1	10000	50
	Production Example 2	SG-2	3500	50
10	Production Example 3	SG-3	9000	50

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TABLE 3

Additives formed in Comparative Production Examples of defoaming agents

		Additive	Weight-average molecular weight	Non-volatile component (%)
20	Comparative Produc- tion Example 1	DN-1	20000	30
	Comparative Production Example 2	DN-2	30000	30
25	Comparative Production Example 3	DN-3	24000	30
	Comparative Production Example 3	DN-4	19000	30
	Comparative Production Example 3	DN-5	22000	30

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TABLE 4

Additives formed in Comparative Production Examples  
of flow-and-leveling agents

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	Additive	Number-average molecular weight	Non-volatile component (%)
Comparative Production Example 1	SN-1	9500	50
Comparative Production Example 2	SN-2	3000	50
Comparative Production Example 3	SN-3	10500	50
Comparative Production Example 4	SN-4	3500	50
Comparative Production Example 5	SN-5	4000	50

Example 1 (Anti-whitening property test of the defoaming agents as  
added to acid anhydride-curing type clear paint, and defoaming test  
in melamine resin-cured clear paint)

Anti-whitening property of painted films was conducted  
with those baked finish type acrylic acid anhydride-curing type clear  
paints of the compositions as shown in Table 5. The same additives  
were also added to those melamine resin-cured clear paint composi-  
tions of Table 6 and their defoaming and anti-popping test was con-  
ducted.

Those starting materials as shown in Table 5 were homoge-  
neously mixed with a dissolver to form an acid anhydride-curing type  
clear paints. To each of the paints 1% by weight to the clear paint of  
one of those defoaming agents shown in Tables 1 and 3 was added and  
dispersed with said dissolver at 2,000 r.p.m. for 2 minutes. Thus  
formed paint formulation each was applied on a glass plate with a 150  
 $\mu$ m-applicator for an anti-whitening property test, and cured by



baking in a 200°C oven for 30 minutes. This painted plate was cooled off to room temperature, immersed in 80°C warm water for an hour. The water was then allowed to cool off to 25°C. The painted plate was withdrawn from the water tank, and water drops on the paint film surface were wiped off with dry non-woven fabric, followed by 24 hours' drying at room temperature. The whitening condition of the paint film was visually observed.

The defoaming test was conducted in the following manner. Viscosity of those clear paints shown in Table 6 was adjusted with a diluting solvent to 25 seconds with Ford cup #4, and each of the paints was applied onto a tin plate with an air spray gun at increasing rates so that the dry paint film thickness should vary from 20 µm to 100 µm. The painted film was allowed to stand for 3 minutes after the application, and then baked in a 160°C oven for 20 minutes to be cured.

The test results of the paint films were evaluated as follows. The anti-whitening property of the baked films formed on the glass plates was visually evaluated in five scales from "the best" (5) to the worst (1). The defoaming property was evaluated by measuring thickness of the dry film, in which foams appeared, on the painted surface of said tin plate with a thickness gauge. Furthermore, the whole painted surface was visually observed, and graded in five scales from "the best" (5) showing the least number of foams to "the worst" (1) showing the greatest number of foams. The test results were as shown in Table 7.

TABLE 5

## Acid Anhydride-Curing Type Clear Paint Formulations

5	Starting Material	Amount (parts)	Maker company
	Finedic A-207S <sup>*1)</sup> (50% solution <sup>*2)</sup>	68.4	Dainippon Ink & Chemicals, Inc.
	Dodecenylsuccinic acid anhydride (50% solution <sup>*2)</sup>	31.6	Wako Pure Chemical Industries, Ltd.
10	DNP-30 (2,4,6-tris(dimethylaminomethyl)phenol)	1.0	Wako Pure Chemical Industries, Ltd.

\*1) epoxy equivalent 490 g/eq.

\*2) diluting solvent: Solvesso #100/xylene/butyl  
cellosolve/n-butanol = 4/3/2/1

TABLE 6

## Melamine Resin-Curing Type Clear Paint

20	Starting Material	Amount (parts)	Maker company
	Beckosol EZ-3530-80	57.9	Dainippon Ink & Chemicals, Inc.
	Super Beckamine L-116-70	28.4	Dainippon Ink & Chemicals, Inc.
	Diluting solvent <sup>*)</sup>	proper quantity	

\*) diluting solvent: Solvesso #100/xylene/butyl  
cellosolve/n-butanol = 4/3/2/1

TABLE 7

## Test Results

Additive	Added amount (%)	Anti-whitening property	Foamed film thickness ( $\mu\text{m}$ )	Number of foams
Blank	—	5	20	1
DG-1	1.0	5	65	3
DG-2	1.0	5	80	5
DG-3	1.0	5	50	3
DN-1	1.0	2	80	5
DN-2	1.0	1	80	5
DN-3	1.0	3	40	2
DN-4	1.0	2	70	4
DN-5	1.0	5	20	1

Example 2 (Anti-whitening property test of the flow-and-leveling agents as added to clear powder coating)

The flow-and-leveling agents were added to the acrylic clear powder coating compositions as shown in Table 8, and the anti-whitening property of the baked films was tested.

[Preparation of master batch]

An acrylic resin (Finedic A-253) was melted at 170°C, in which each one of those flow-and-leveling agents of Tables 2 and 4 was uniformly dispersed to the solid concentration of 10%, with a high speed dissolver. The dispersions were cooled, pulverized with a pin-mill and sieved through a 32 mesh-sieve to provide 10% acrylic resin master batch compositions.

[Preparation of acrylic powder coating]

Acrylic powder coatings of Table 8, which contained the above master batch compositions, were dry-blended and then melt-

kneaded in an extruder maintained at 100-110°C. The kneaded masses were cooled, pulverized in a pin-mill, and sieved through a 150 mesh-sieve to provide white, clear powder coatings.

[Application of the acrylic powder coatings and evaluation]

- 5                Each of above powder coatings was blown onto a glass plate by electrostatic powder coating method, to the cured film thickness of 100  $\mu\text{m}$ , and the anti-whitening property of the films was evaluated. The powder coatings were also blown onto a thermoresistant stainless steel sheet (0.5 x 70 x 100 mm) each, by electrostatic powder coating
- 10 method, to the cured film thickness of 40  $\mu\text{m}$ . Those coated glass plates and stainless steel sheets were then baked at 160°C for 30 minutes. Evaluation of anti-whitening property was conducted in the same manner with Example 1, and the flow-and-leveling property of the baked films was visually evaluated. The results were as shown in
- 15 Table 9.

TABLE 8

Acrylic Clear Coating Powder

Starting Material	Amount (parts)	Maker Company
Acrylic resin : Finedic A-253	75.5	Dainippon Ink & Chemicals, Inc.
Curing agent : Dodecandioic acid	21	Wako Pure Chemical Industries, Ltd.
Flow-and-leveling agent: acrylic resin master batch	5	

TABLE 9

Flow-and-Leveling Property Test Results  
of Acrylic Clear Powder Coatings

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Additive	Amount (%)	Anti-whitening property	Flow-and-leveling property
Blank	—	5	1
SG-1	0.5	5	4
SG-2	0.5	5	5
SG-3	0.5	5	3
SN-1	0.5	1	4
SN-2	0.5	3	5
SN-3	0.5	1	3
SN-4	0.5	2	4
SN-5	0.5	5	1

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